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(54) Title: USE OF FACTOR X POLYMORPHISM IN THE DIAGNOSIS AND TREATMENT OF FACTOR X AND/OR FACTOR XA MEDIATED DISEASES

(57) Abstract

This invention relates to polymorphisms in the human Factor X gene, in particular to the discovery of two single nucleotide polymorphisms in the coding sequence of the human Factor X gene. The invention also relates to methods and materials for analysing allelic variation in the Factor X gene, and to the use of Factor X polymorphism in the diagnosis and treatment of Factor X and/or Factor Xa–mediated diseases, such as thrombotic diseases.